Identification of genomic regions carrying a causal mutation in unordered genomes

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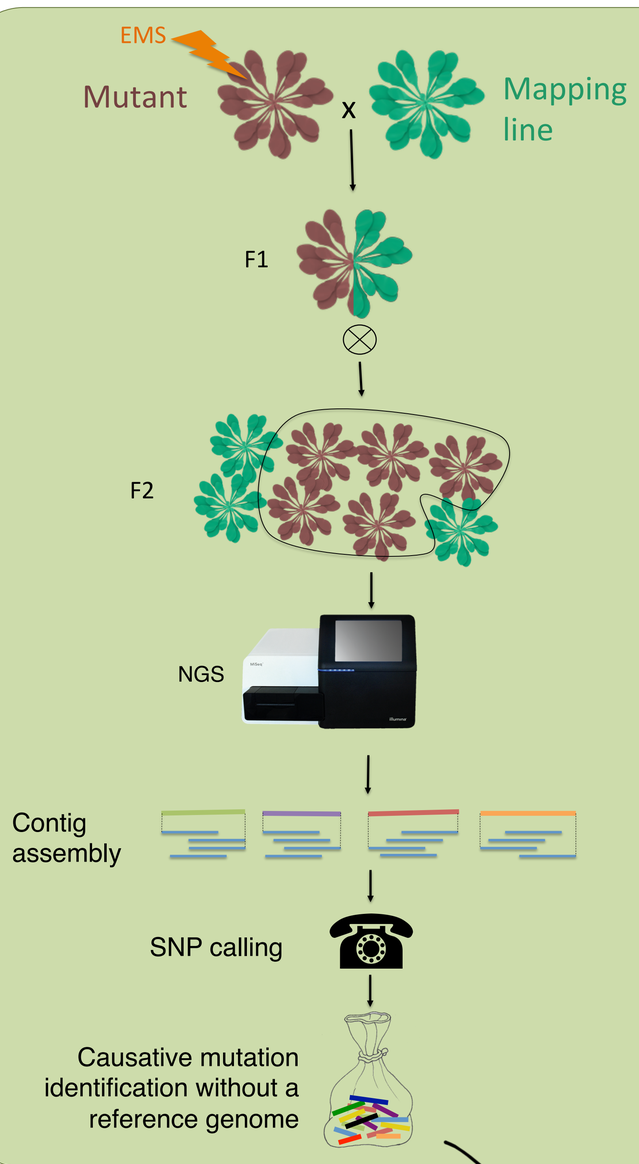


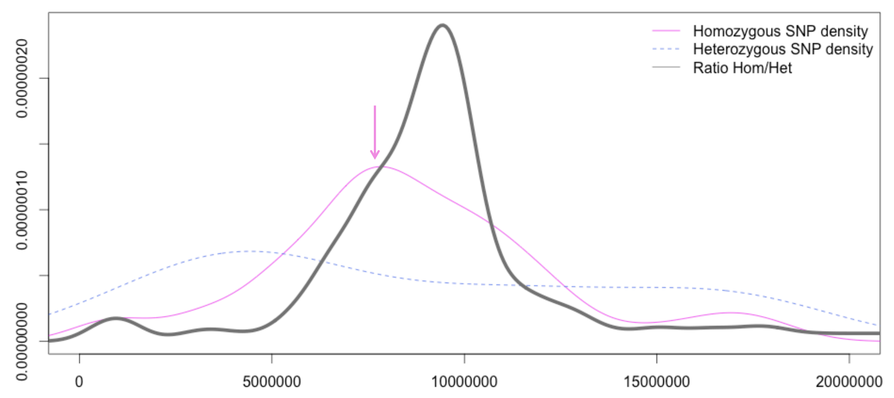
Motivation

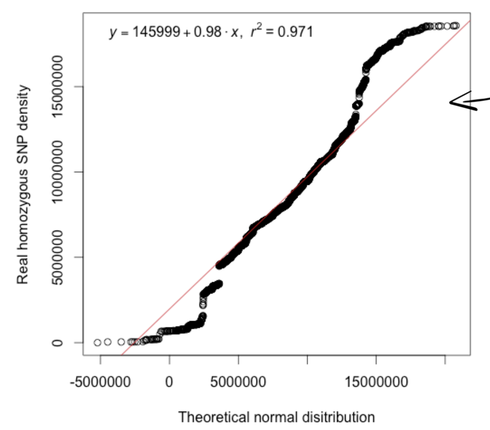
Forward genetic screens have been a fundamental strategy to find genes involved in biological pathways in model species. Mutagenized individuals with a phenotype of interest are isolated and a recombinant mapping population is created by back-crossing to the parental line or out-crossing to a polymorphic ecotype.

The recombination frequency between the causal mutation and nearby genetic markers is low, so the alleles of these genetic markers will co-segregate with the phenotype-altering mutation while the remaining unlinked makers segregate randomly in the genome. Hence, the allele distribution analysis can unhide these low recombinant regions to identify the location of the causal mutation.

Traditional genetic mapping is a work intensive and time-consuming process but recent advances in high-throughput sequencing (HTS) have greatly accelerated the identification of mutations underlying mutant phenotypes in forward genetic screens. In the last few years, researchers have developed user-friendly tools for mapping-by-sequencing, yet they are not applicable to **organisms with non-sequenced genomes.**

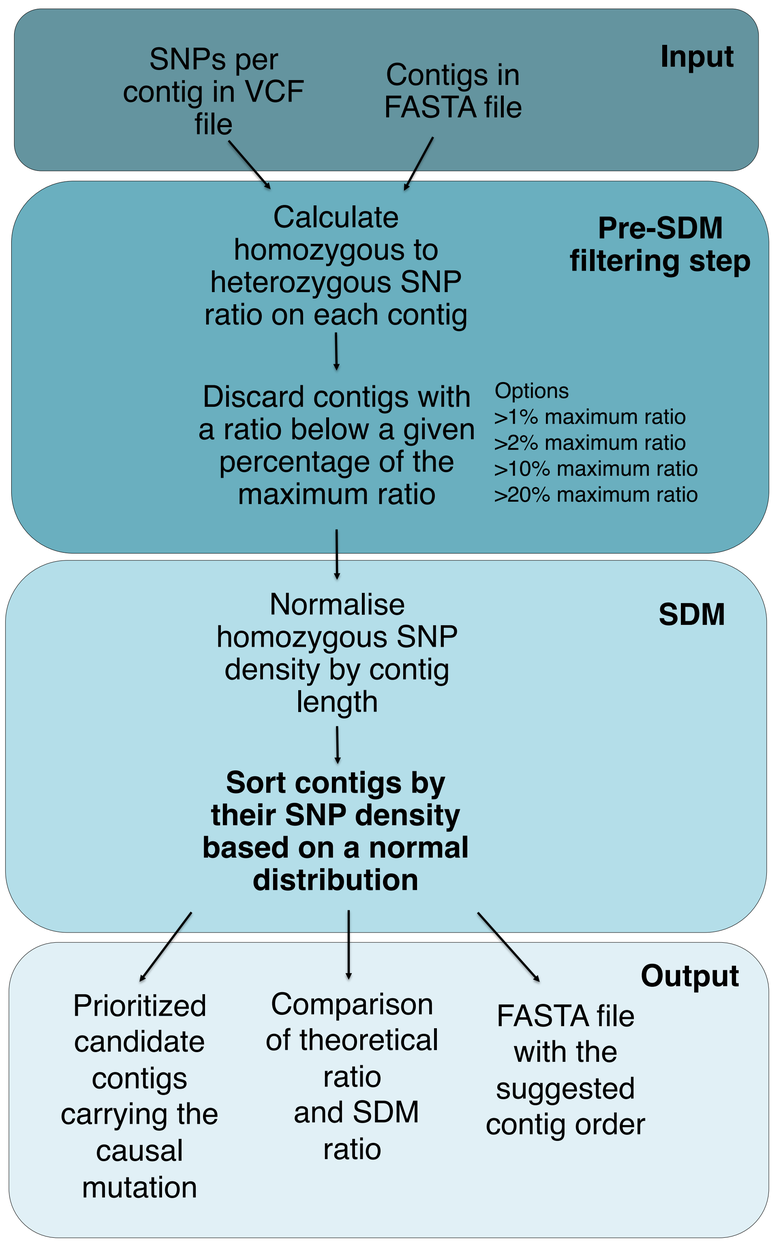
SNP density plots revealed the homozygous SNP linkage around the causative mutation causing a high homozygous to heterozygous ratio signal where the mutation is located



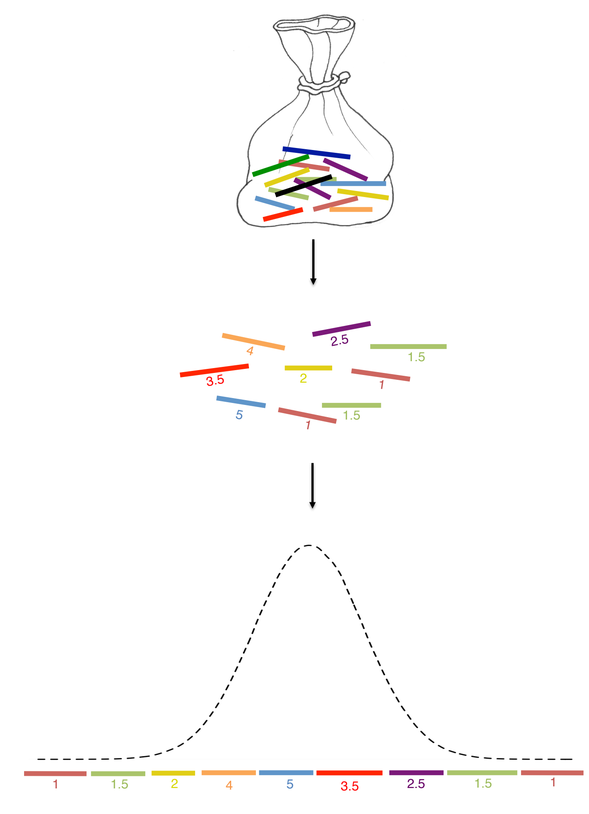


Homozygous SNPs are normally distributed around the causal mutation!

SNP Distribution Method

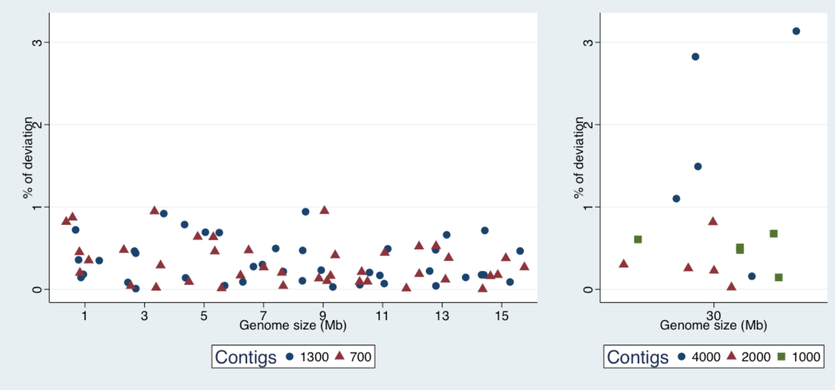


SDM is a fast causative mutant identification method based on a simple reference-free contig assembly that allows the detection of candidate causative SNPs. Instead of relying on a genome comparison, it focus on the SNP linkage around the causal mutation and analyse the SNP distribution to identify the chromosome area where the putative mutated gene is located.



Modelling

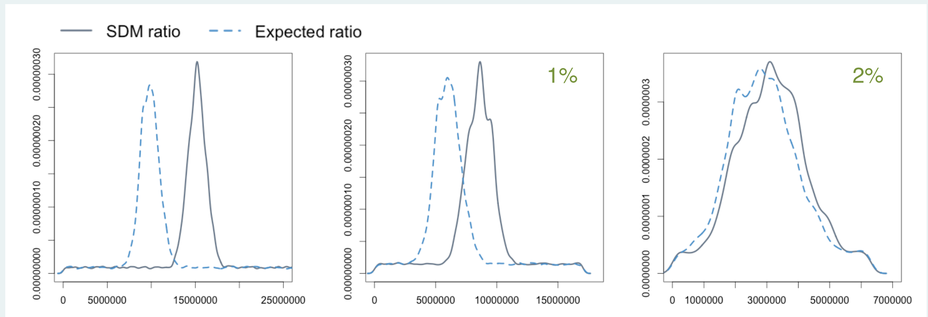
Model genomes are useful to help us developing our method and identifying its limitations. By using idealised SNP distributions, we can predict were the mutation is going to be located and estimate the deviation of SDM from this expected position. A normal distribution was used for the homozygous SNPs while heterozygous SNPs followed a uniform distribution. We created different model genomes based on *Arabidopsis thaliana* chromosome I. We tested the effect of genome length and contig size on SDM performance.



We define the homozygous to heterozygous SNP ratio on contig n as:

$$Ratio_n = \frac{(\sum Hom) + 1}{(\sum Het) + 1}$$

The contigs that are located far from the causal mutation have a constant homozygous SNP density due to recombination. The low ratio in these regions is used as a filter to focus on the genomic region where the mutation is likely to be found. Contigs with a ratio falling below a % of the maximum ratio will be discarded.



Take home

- ✓ **Forward genetic screens** are very useful to identify genes responsible for particular phenotypes.
- ✓ Homozygous SNPs are **normally distributed** in the mutant genome of back-cross and out-crossed individuals. We defined a theoretical SNP distribution used by SDM to identify the genomic region where the causative mutation is located.
- ✓ SDM does not rely on previously known genetic markers and can be used on extremely **fragmentary genome assemblies**, even down to the level of long reads.

I'm a predoctoral intern at The Sainsbury Laboratory doing Bioiformatics in the team MacLean.

/pilarcormo/
SNP_Distribution_method

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